```
=> file biosis caba caplus embase japio lifesci medline scisearch
=> e bisen prakash/au
E1
            7
                  BISEN P S */AU
E2
            4
                  BISEN P S DR/AU
E3
            2 --> BISEN PRAKASH/AU
E4
           124
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E5
            1
                  BISEN PRAKASH S PROF/AU
E6
           32
                  BISEN PRAKASH SINGH/AU
E7
           33
                  BISEN PS/AU
Ε8
            1
                  BISEN R/AU
E9
           24
                  BISEN R K/AU
E10
           12
                  BISEN R S/AU
E11
            1
                 BISEN RUTH/AU
E12
            1
                 BISEN S/AU
=> s e1-e7 and tuberculosis and diagnos?
            20 ("BISEN P S *"/AU OR "BISEN P S DR"/AU OR "BISEN PRAKASH"/AU OR
               "BISEN PRAKASH S"/AU OR "BISEN PRAKASH S PROF"/AU OR "BISEN PRAK
              ASH SINGH"/AU OR "BISEN PS"/AU) AND TUBERCULOSIS AND DIAGNOS?
=> dup rem 11
PROCESSING COMPLETED FOR L1
             7 DUP REM L1 (13 DUPLICATES REMOVED)
=> s 12 and glycolipid?
             2 L2 AND GLYCOLIPID?
L3
=> d bib ab kwic 1-
YOU HAVE REQUESTED DATA FROM 2 ANSWERS - CONTINUE? Y/(N):y
     ANSWER 1 OF 2 BIOSIS COPYRIGHT (c) 2010 The Thomson Corporation on STN
AN
     2005:464785 BIOSIS <<LOGINID::20100824>>
    PREV200510248351
DN
      ***Glycolipids*** of Mycobacterium ***tuberculosis***
ΤI
                                                                  strain H37Rv
     are potential serological markers for ***diagnosis*** of active
      ***tuberculosis***
     Tiwari, R. P.; Tiwari, Dileep; Garq, Sanjay K.; Chandra, Ramesh;
ΑU
       ***Bisen, Prakash S.*** [Reprint Author]
     Bundelkhand Univ, JC Bose Inst Life Sci, Dept Biotechnol, Jhansi 284218,
CS
     Uttar Pradesh, India
     prakash bisen@hotmail.com
    Clinical and Diagnostic Laboratory Immunology, (MAR 2005) Vol. 12, No. 3,
SO
     pp. 465-473.
     ISSN: 1071-412X.
DT
    Article
LA
    English
     Entered STN: 9 Nov 2005
ΕD
     Last Updated on STN: 9 Nov 2005
     A simple and cost-effective ***diagnostic*** tool (TB Screen Test) for
AΒ
     the screening of patients with pulmonary and extrapulmonary
       ***tuberculosis*** and for differentiation of those individuals from
     individuals without ***tuberculosis*** , other common infections, and
     healthy controls has been developed. The serological responses of
     purified mycobacterial ***qlycolipid*** antigens were examined by a
     liposome agglutination assay. The assay was able to detect very low
     antiglycolipid antibody concentrations in the infected individuals. The
     sera from the ***tuberculosis*** patient group had significantly
```

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higher concentrations of antiglycolipid antibody than the sera from
     uninfected control subjects, with 94% sensitivity and 98.3% specificity.
       ***Glycolipids*** of Mycobacterium ***tuberculosis*** H37Rv
antigens
    were isolated, purified, and characterized. After interchelation with
     liposome particles, these purified antigens specifically bound to the
     antiglycolipid antibodies present in the sera of patients with
      ***tuberculosis*** , resulting in the formation of a blue agglutination.
     This protocol clearly differentiates healthy controls and M. bovis
     BCG-vaccinated subjects from those with active ***tuberculosis***
                   ***diagnostic*** tool, the TB Screen Test, is more
     The resultant
     economical and rapid (4 min) than other currently available products and
     can be used for the mass screening of a heavily afflicted population.
ΤI
      ***Glycolipids*** of Mycobacterium ***tuberculosis*** strain H37Rv
     are potential serological markers for ***diagnosis*** of active
      ***tuberculosis***
ΑU
    Tiwari, R. P.; Tiwari, Dileep; Garg, Sanjay K.; Chandra, Ramesh;
       ***Bisen, Prakash S.*** [Reprint Author]
    A simple and cost-effective ***diagnostic*** tool (TB Screen Test) for
AΒ
     the screening of patients with pulmonary and extrapulmonary
       ***tuberculosis*** and for differentiation of those individuals from
     individuals without ***tuberculosis*** , other common infections, and
     healthy controls has been developed. The serological responses of
     purified mycobacterial ***glycolipid*** antigens were examined by a
     liposome agglutination assay. The assay was able to detect very low
     antiglycolipid antibody concentrations in the infected individuals. The
                  ***tuberculosis*** patient group had significantly
     higher concentrations of antiglycolipid antibody than the sera from
    uninfected control subjects, with 94% sensitivity and 98.3% specificity.
      ***Glycolipids*** of Mycobacterium ***tuberculosis*** H37Rv
antigens
     were isolated, purified, and characterized. After interchelation with
     liposome particles, these purified antigens specifically bound to the
     antiglycolipid antibodies present in the sera of patients with
      ***tuberculosis*** , resulting in the formation of a blue agglutination.
     This protocol clearly differentiates healthy controls and M. bovis
     BCG-vaccinated subjects from those with active ***tuberculosis***
     The resultant ***diagnostic*** tool, the TB Screen Test, is more
     economical and rapid (4 min) than other currently available products and
     can be. . .
    Major Concepts
       Infection; Clinical Chemistry (Allied Medical Sciences)
ΤТ
    Diseases
            ***tuberculosis*** : bacterial disease, ***diagnosis***
           ***Tuberculosis***
                               (MeSH)
ΙT
    Chemicals & Biochemicals
           ***glycolipids*** ; serological markers
IT
    Methods & Equipment
       TB Screen Test: clinical techniques, ***diagnostic*** techniques
ORGN Classifier
       Mycobacteriaceae
                         08881
     Super Taxa
       Mycobacteria; Actinomycetes and Related Organisms; Eubacteria;
       Bacteria; Microorganisms
     Organism Name
       Mycobacterium bovis (species): pathogen
       Mycobacterium ***tuberculosis*** (species): pathogen, strain-H37Rv
```

Taxa Notes

Bacteria, Eubacteria, Microorganisms

- L3 ANSWER 2 OF 2 EMBASE COPYRIGHT (c) 2010 Elsevier B.V. All rights reserved on STN
- AN 2007510106 EMBASE <<LOGINID::20100824>>
- TI Rapid liposomal agglutination card test for the detection of antigens in patients with active ***tuberculosis****.
- AU Tiwari, R.P.
- CS Diagnostic Division, Nicholas Piramal India Limited, Pawane, Navi, Mumbai, India.
- AU Tiwari, R.P.; Garg, S.K.; ***Bisen, Prakash S. (correspondence)***
- CS Institute of Biotechnology and Allied Sciences, Seedling Academy of Design, Technology and Management, Jagatpura, Jaipur, India. psbisen@gmail.com
- AU Garg, S.K.
- CS Department of Biochemistry, University of Nebraska, Lincoln, NE, United States.
- AU Bharmal, R.N.; Kartikeyan, S.
- CS Department of Microbiology, Preventive and Social Medicine, Rajiv Gandhi Medical College, Kalwa, Thane, India.
- AU ***Bisen, Prakash S. (correspondence) ***
- CS Bisen Biotech and Biopharma Pvt. Ltd., M-7 Laxmipuram, Transport Nagar, Gwalior 474009, India. psbisen@gmail.com
- SO International Journal of Tuberculosis and Lung Disease, (Oct 2007) Vol. 11, No. 10, pp. 1143-1151.

Refs: 30

ISSN: 1027-3719 CODEN: IJTDFO

- CY France
- DT Journal; Article
- FS 015 Chest Diseases, Thoracic Surgery and Tuberculosis
 004 Microbiology: Bacteriology, Mycology, Parasitology and Virology
 006 Internal Medicine
- LA English
- SL English; French; Spanish; Castilian
- ED Entered STN: 30 Oct 2007
 Last Updated on STN: 30 Oct 2007
- SETTING: A total of 1360 subjects with clinically confirmed pulmonary and AΒ extra-pulmonary ***tuberculosis*** (TB) and other non-tuberculous conditions. OBJECTIVES: To develop a rapid, sensitive and specific ***diagnostic*** test for the detection of the ***glycolipid*** ***tuberculosis*** in a variety of clinical antigen of Mycobacterium samples. STUDY DESIGN: Affinity-purified rabbit anti- ***glycolipid*** antibodies (IgG) were coupled to liposome particles (0.2-0.4 .mu.m) in the presence of 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride and N-hydroxysuccinamide to prepare the working reagent of the TB/M card test. RESULTS: Antibody-conjugated liposomes, when determined with the ***glycolipid*** antigens present in the specimens, formed a dark blue agglutination within 4 min. No dumping was observed in samples from normal healthy subjects or patients with other diseases. The test was shown to be effective in detecting ***glycolipid*** antigens of M.

with

as low as 1 ng/ml analytical sensitivity, 97.4% clinical sensitivity and 96.9% specificity. CONCLUSION: The TB/M card test was found to be comparatively economical (4 Indian Rupees or US\$ 0.09/test), rapid (4 min) and seems fairly useful for mass testing of a variety of biological

tuberculosis in clinical samples from patients with active TB

```
specimens (cerebrospinal, pleural and synovial fluids, serum, tissue
     biopsy extract) from patients with tuberculous meningitis, pulmonary TB
     and other extra-pulmonary TB in endemic countries. . COPYRGT. 2007 The
     Union.
     Rapid liposomal agglutination card test for the detection of antigens in
     patients with active ***tuberculosis*** .
     Tiwari, R.P.; Garg, S.K.; ***Bisen, Prakash S. (correspondence)***
     Institute of Biotechnology and Allied Sciences, Seedling Academy of
     Design, Technology and Management, Jagatpura, Jaipur, India
       ***Bisen, Prakash S. (correspondence) ***
     Bisen Biotech and Biopharma Pvt. Ltd., M-7 Laxmipuram, Transport Nagar,
     Gwalior 474009, India. psbisen@gmail.com
     SETTING: A total of 1360 subjects with clinically confirmed pulmonary and
     extra-pulmonary
                      ***tuberculosis*** (TB) and other non-tuberculous
     conditions. OBJECTIVES: To develop a rapid, sensitive and specific
       ***diagnostic*** test for the detection of the
                                                        ***glycolipid***
     antigen of Mycobacterium ***tuberculosis***
                                                   in a variety of clinical
     samples. STUDY DESIGN: Affinity-purified rabbit anti- ***qlycolipid***
     antibodies (IgG) were coupled to liposome particles (0.2-0.4 .mu.m) in the
     presence of 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride
     and N-hydroxysuccinamide to prepare the working reagent of the TB/M card
     test. RESULTS: Antibody-conjugated liposomes, when determined with the
      ***qlycolipid*** antigens present in the specimens, formed a dark blue
     agglutination within 4 min. No dumping was observed in samples from
     normal healthy subjects or patients with other diseases. The test was
     shown to be effective in detecting ***qlycolipid*** antigens of M.
       ***tuberculosis*** in clinical samples from patients with active TB
with
     as low as 1 ng/ml analytical sensitivity, 97.4% clinical sensitivity and.
    Medical Descriptors:
     adolescent
     adult
     *agglutination test
     *antigen detection
     article
     cerebrospinal fluid
     controlled study
         ***diagnostic test***
         ***extrapulmonary tuberculosis***
     human
         ***lung tuberculosis***
     major clinical study
         ***Mycobacterium tuberculosis***
     pleura fluid
     priority journal
     school child
     sensitivity and specificity
     synovial fluid
         ****tuberculosis***
     tuberculous meningitis
     1 (3 dimethylaminopropyl) 3 ethylcarbodiimide
     amide
     antibody conjugate
         ***qlycolipid***
     liposome
```

ТΤ

ΑIJ

CS

ΑU

CS

AΒ

n hydroxysuccinamide

```
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E1
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                  TIWARI RAKESH VALLABHDAS/AU
E2
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                  TIWARI RAKSHA/AU
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E.4
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E11
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ΑN
    PREV200400038179
DΝ
TΙ
    Analysis of the shotgun expression library of the Mycobacterium
      ***tuberculosis*** genome for immunodominant polypeptides: Potential
use
    in serodiagnosis.
    Bisen, Prakash S. [Reprint Author]; Garq, Sanjay K.; ***Tiwari, Ram***
        P.***; Tagore, P. Ravindra Nath; Chandra, Ramesh; Karnik, Rucha;
Thaker,
    Nimesh; Desai, Nirav; Ghosh, P. K.; Fraziano, Maurizio; Colizzi, Vittorio
    Madhav Institute of Technology and Science, Gwalior, MP, 474 005, India
CS
     prakash_bisen@hotmail.com
    Clinical and Diagnostic Laboratory Immunology, (November 2003) Vol. 10,
    No. 6, pp. 1051-1058. print.
     ISSN: 1071-412X (ISSN print).
DT
    Article
LA
    English
    Entered STN: 7 Jan 2004
ED
     Last Updated on STN: 7 Jan 2004
=> s 15 and glycolipid?
            0 L5 AND GLYCOLIPID?
=> e tiwari ram pramod/au
```

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TIWARI RAM PRAKAH/AU
E1
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E2
            20
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Е3
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L8
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     2007:385701 BIOSIS <<LOGINID::20100824>>
ΑN
    PREV200700390349
DN
    Modern approaches to a rapid diagnosis of tuberculosis: Promises and
     challenges ahead.
       ***Tiwari, Ram Pramod*** ; Hattikudur, Narendra S.; Bharmal, Ramesh N.;
ΑU
     Kartikeyan, S.; Deshmukh, Neeta M.; Bisen, Prakash S. [Reprint Author]
     Seeding Acad Design Technol and Management, Inst Biotechnol and Allied
     Sci, Jaipur 302004, Rajasthan, India
    psbisen@gmail.com
SO
    Tuberculosis (Amsterdam), (MAY 2007) Vol. 87, No. 3, pp. 193-201.
    ISSN: 1472-9792.
DT
    Article
    General Review; (Literature Review)
LA
    English
ED
    Entered STN: 11 Jul 2007
     Last Updated on STN: 11 Jul 2007
=> s liposom? and tuberculosis and glycolipid and antibod?
L9
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=> dup rem 19
PROCESSING COMPLETED FOR L9
L10
            10 DUP REM L9 (13 DUPLICATES REMOVED)
=> s 110 and diagnos?
            5 L10 AND DIAGNOS?
=> d bib ab kwic 1-
YOU HAVE REQUESTED DATA FROM 5 ANSWERS - CONTINUE? Y/(N):y
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L11 ANSWER 1 OF 5 BIOSIS COPYRIGHT (c) 2010 The Thomson Corporation on STN

- AN 2005:464785 BIOSIS <<LOGINID::20100824>>
- DN PREV200510248351
- TI Glycolipids of Mycobacterium ***tuberculosis*** strain H37Rv are potential serological markers for ***diagnosis*** of active ***tuberculosis***.
- AU Tiwari, R. P.; Tiwari, Dileep; Garg, Sanjay K.; Chandra, Ramesh; Bisen, Prakash S. [Reprint Author]
- CS Bundelkhand Univ, JC Bose Inst Life Sci, Dept Biotechnol, Jhansi 284218, Uttar Pradesh, India prakash bisen@hotmail.com
- SO Clinical and Diagnostic Laboratory Immunology, (MAR 2005) Vol. 12, No. 3, pp. 465-473.
 ISSN: 1071-412X.
- DT Article
- LA English
- ED Entered STN: 9 Nov 2005 Last Updated on STN: 9 Nov 2005
- AΒ A simple and cost-effective ***diagnostic*** tool (TB Screen Test) for the screening of patients with pulmonary and extrapulmonary ***tuberculosis*** and for differentiation of those individuals from individuals without ***tuberculosis*** , other common infections, and healthy controls has been developed. The serological responses of purified mycobacterial ***glycolipid*** antigens were examined by a ***liposome*** agglutination assay. The assay was able to detect very low antiglycolipid ***antibody*** concentrations in the infected individuals. The sera from the ***tuberculosis*** patient group had significantly higher concentrations of antiglycolipid ***antibody*** than the sera from uninfected control subjects, with 94% sensitivity and 98.3% specificity. Glycolipids of Mycobacterium ***tuberculosis*** H37Rv antigens were isolated, purified, and characterized. After interchelation with ***liposome*** particles, these purified antigens specifically bound to the antiglycolipid ***antibodies*** present in the sera of patients with ***tuberculosis*** , resulting in the formation of a blue agglutination. This protocol clearly differentiates healthy controls and M. bovis BCG-vaccinated subjects from those with active ***tuberculosis*** . The resultant ***diagnostic*** tool, the TB Screen Test, is more economical and rapid (4 min) than other currently available products and can be used for the mass screening of a heavily afflicted population.
- TI Glycolipids of Mycobacterium ***tuberculosis*** strain H37Rv are potential serological markers for ***diagnosis*** of active ***tuberculosis*** .
- AB A simple and cost-effective ***diagnostic*** tool (TB Screen Test) for the screening of patients with pulmonary and extrapulmonary ***tuberculosis*** and for differentiation of those individuals from individuals without ***tuberculosis***, other common infections, and healthy controls has been developed. The serological responses of purified mycobacterial ***glycolipid*** antigens were examined by a ***liposome*** agglutination assay. The assay was able to detect very low antiglycolipid ***antibody*** concentrations in the infected individuals. The sera from the ***tuberculosis*** patient group had significantly higher concentrations of antiglycolipid ***antibody*** than the sera from uninfected control subjects, with 94% sensitivity and 98.3% specificity. Glycolipids of Mycobacterium ***tuberculosis*** H37Rv antigens were isolated, purified, and characterized. After interchelation with ***liposome*** particles, these purified antigens specifically bound to the antiglycolipid ***antibodies*** present in

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IT Major Concepts

Infection; Clinical Chemistry (Allied Medical Sciences)

IT Diseases

tuberculosis : bacterial disease, ***diagnosis***
Tuberculosis (MeSH)

IT Chemicals & Biochemicals

glycolipids; serological markers

IT Methods & Equipment

TB Screen Test: clinical techniques, ***diagnostic*** techniques ORGN Classifier

Mycobacteriaceae 08881

Super Taxa

Mycobacteria; Actinomycetes and Related Organisms; Eubacteria; Bacteria; Microorganisms

Organism Name

Mycobacterium bovis (species): pathogen

Mycobacterium ***tuberculosis*** (species): pathogen, strain-H37Rv Taxa Notes

Bacteria, Eubacteria, Microorganisms

L11 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2010 ACS on STN

AN 1997:425363 CAPLUS <<LOGINID::20100824>>

DN 127:32828

OREF 127:6345a,6348a

- TI Therapeutic and ***diagnostic*** vaccine for the treatment of microbial infections
- IN Pascual, David; Bond, Clifford; Burritt, James; Burgess, Don; Glee, Pati;
 Jutila, John; Jutila, Mark; Bargatze, Robert; Mcfeters, Gordon; Pyle,
 Barry; Cutler, Jim E.; Han, Yongmoon
- PA Research and Development Institute, Inc., USA; Pascual, David; Bond, Clifford; Burritt, James; Burgess, Don; Glee, Pati; Jutila, John; Jutila, Mark; Bargatze, Robert; et al.
- SO PCT Int. Appl., 98 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 4

	PATENT NO.					KIND		DATE		APPLICATION NO.						DATE			
PI		9718790 9718790						 19970529 19970731			WO 1996-US18796				19961121				
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			EE,	ES,	FI,	GB,	GE,	HU,	IL,	IS,	JP,	KE,	KG,	KP,	KR,	KΖ,	LC,	LK,	
			LR,	LS,	LT,	LU,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	NΖ,	PL,	PT,	RO,	RU,	
			SD,	SE,	SG,	SI,	SK,	ΤJ,	TM,	TR,	TT,	UA,	UG,	US,	UZ,	VN			
		RW:	ΚE,	LS,	MW,	SD,	SZ,	UG,	AT,	BE,	CH,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	
			ΙE,	ΙΤ,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	$\mathrm{ML}_{m{\prime}}$	
			MR,	NE,	SN,	TD,	ΤG												
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	AU	9711226				Α		19970611			AU 1997-11226					19961121			
	EP	IP 869801			A2		19981014			EP 1996-942049					19961121				

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EP 869801
                               20040121
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        R: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LI, LU, MC, NL,
            PT, SE
                         Τ
                               20000328
                                        JP 1997-519932
    JP 2000503630
                                                                 19961121
    AT 258057
                        Τ
                               20040215 AT 1996-942049
                                                                19961121
                     A1 20041209 US 2004-780650
    US 20040247611
                                                                20040219
PRAI US 1995-7477P
                       Р
                             19951122
    US 1994-247972
                       B2 19940523
                       A2
    US 1995-483558
                              19950607
    WO 1996-US18796
                        W
                               19961121
    US 1998-68935
                        В1
                               19981123
    Therapeutic peptides, vaccines and
                                        ***diagnostic*** agents are
AB
    disclosed for the treatment of pathogenic infections. The agents are
     capable of binding to mol. address on host cell (e.g. leukocyte,
     endothelial or epithelial cells, nerve cells), triggering one or more
     signal transduction pathways and enabling selective pathogen or toxin to
     traffic through host tissue. The agents are microbial attachment mols.
     such as adhesive protein, glycoprotein, lectin, carbohydrate,
      ***glycolipid***
             THERE ARE 4 CAPLUS RECORDS THAT CITE THIS RECORD (4 CITINGS)
OSC.G
RE.CNT 3
             THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
ΤI
     Therapeutic and ***diagnostic*** vaccine for the treatment of
    microbial infections
AB
    Therapeutic peptides, vaccines and ***diagnostic*** agents are
    disclosed for the treatment of pathogenic infections. The agents are
     capable of binding to mol. address on host. . . or toxin to traffic
     through host tissue. The agents are microbial attachment mols. such as
     adhesive protein, glycoprotein, lectin, carbohydrate,
                                                          ***qlvcolipid***
ST
    microbial adhesion mol vaccine
                                   ***diagnostic*** ; monoclonal
      ***antibody*** microbial antigen therapeutic
                                                      ***diagnostic***
    Agglutinins and Lectins
ΤТ
    RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
       (C-type (calcium-dependent type); vaccine comprising microbial adhesion
       mol. antigen as therapeutic and ***diagnostic*** for microbial
       infections)
ΙT
    Integrins
    RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
       (CD41a; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΤТ
    Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (CD49; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
    Gene, animal
ΙT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (Cdc42, CTP-binding protein; vaccine comprising microbial adhesion mol.
       antigen as therapeutic and ***diagnostic*** for microbial
       infections)
ΙT
    Selectins
    RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
       (E-; vaccine comprising microbial adhesion mol. antigen as therapeutic
       and ***diagnostic*** for microbial infections)
    Proteins, specific or class
ΙT
```

```
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (GTP-binding; vaccine comprising microbial adhesion mol. antigen as
                     ***diagnostic*** for microbial infections)
   therapeutic and
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (ICAM-1 (intercellular adhesion mol. 1); vaccine comprising microbial
   adhesion mol. antigen as therapeutic and ***diagnostic***
   microbial infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (ICAM-2 (intercellular adhesion mol. 2); vaccine comprising microbial
   adhesion mol. antigen as therapeutic and ***diagnostic***
   microbial infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (ICAM-3 (intercellular adhesion mol. 3); vaccine comprising microbial
   adhesion mol. antigen as therapeutic and ***diagnostic***
  microbial infections)
Selectins
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
   (L-; vaccine comprising microbial adhesion mol. antigen as therapeutic
   and ***diagnostic***
                          for microbial infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (MAdCAM-1; vaccine comprising microbial adhesion mol. antigen as
   therapeutic and
                   ***diagnostic*** for microbial infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (N-CAM; vaccine comprising microbial adhesion mol. antigen as
                   ***diagnostic*** for microbial infections)
   therapeutic and
Selectins
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
   (P-; vaccine comprising microbial adhesion mol. antigen as therapeutic
   and ***diagnostic*** for microbial infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (PECAM-1; vaccine comprising microbial adhesion mol. antigen as
   therapeutic and ***diagnostic*** for microbial infections)
Toxins
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (Shiga-like toxin, Escherichia coli; vaccine comprising microbial
   adhesion mol. antigen as therapeutic and ***diagnostic***
  microbial infections)
Antigens
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (Streptococcal SA I/II; vaccine comprising microbial adhesion mol.
   antigen as therapeutic and
                              ***diagnostic*** for microbial
   infections)
Cell adhesion molecules
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (VAM-1; vaccine comprising microbial adhesion mol. antigen as
   therapeutic and ***diagnostic*** for microbial infections)
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RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL

(Biological study); USES (Uses)

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(VLA; vaccine comprising microbial adhesion mol. antigen as therapeutic ***diagnostic*** for microbial infections) and ΙT Intestine (adhesion of Escherichia coli; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΤТ ***Diagnosis*** (agents; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΙT Macrophage (alveolar; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT (alveolus, macrophage; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT Integrins RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (antigens Mac-1 (macrophage 1); vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) Epithelium ΤT Respiratory tract (cells; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΤT Peptides, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (conjugates; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT Blood vessel (endothelium, cells; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΤT Toxins RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (enterotoxins, Escherichia coli; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) Clostridium botulinum ΙT Clostridium tetani (exotoxin; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΤT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (exotoxins; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΤТ G proteins (guanine nucleotide-binding proteins) RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (gene TC4; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ITG proteins (guanine nucleotide-binding proteins) RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (gene rab; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΤТ G proteins (quanine nucleotide-binding proteins) RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (gene rac; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections)

ΙT Integrins RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (gp150.95; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΤТ Digestive tract Digestive tract (hemorrhage, microorganism causing; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** microbial infections) ΙT Urogenital tract (infection; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT Integrins RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (leucam; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT (macrophage; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) Sialic acids ITRL: BSU (Biological study, unclassified); BIOL (Biological study) (microbial adhesion mol. contg.; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) Glycopeptides ΤТ Peptides, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (microbial adhesion mol.; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) Agglutinins and Lectins ΤT Ligands RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (microbial; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ***Antibodies*** ΙT RL: BPN (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses) (monoclonal; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΤТ Pharynx (nasopharynx, epithelium and endothelium; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT Nerve (neuron; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΙT Infection (nosocomial; vaccine comprising microbial adhesion mol. antigen as ***diagnostic*** for microbial infections) therapeutic and ΤT Peptides, biological studies RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (oligopeptides; vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections) ΙT Urinary tract

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(pathogen; vaccine comprising microbial adhesion mol. antigen as
        therapeutic and ***diagnostic*** for microbial infections)
ΙT
        (phycomycetous; vaccine comprising microbial adhesion mol. antigen as
                         ***diagnostic*** for microbial infections)
        therapeutic and
ΙT
    Paramagnetic materials
        (superparamagnetic, beads; vaccine comprising microbial adhesion mol.
        antigen as therapeutic and ***diagnostic*** for microbial
        infections)
ΙT
     Escherichia coli
        (uropathjogenic; vaccine comprising microbial adhesion mol. antigen as
        therapeutic and ***diagnostic*** for microbial infections)
ΙT
    Animal cell
    Animal tissue
     Aspergillus
     B cell (lymphocyte)
     Bacteriophage
     Blastomyces
     Bordetella pertussis
     Brucella
     Candida
     Candida albicans
     Cell adhesion
     Chlamydia
    Coccidioides
     Coliphage M13
     Corynebacterium diphtheriae
     Cowpea mosaic virus
     Cryptococcus (fungus)
     Cryptosporidium
         ***Diagnosis***
     Entamoeba histolytica
     Enterobacter aerogenes
     Eukaryote (Eukaryotae)
     Francisella tularensis
     Fungi
     Genetic vectors
     Giardia lamblia
     Haemophilus influenzae
     Hafnia alvei
     Hantavirus
    Helicobacter pylori
     Hepatitis virus
     Histoplasma
     Human adenovirus
     Human coxsackievirus
     Human herpesvirus
     Human immunodeficiency virus
     Human poliovirus
     Influenza A virus
     Influenza B virus
     Influenza C virus
    Klebsiella pneumoniae
     Legionella
     Leishmania
     Leukocyte
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Liposomes

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Measles virus
Microorganism
Mumps virus
                ***tuberculosis***
Mycobacterium
Mycoplasma pneumoniae
Neisseria gonorrhoeae
Neisseria meningitidis
Organ, animal
Parasite
Pathogen
Pilus
Plasmodium berghei
Plasmodium falciparum
Prokaryote
Proteus (bacterium)
Pseudomonas
Pseudomonas aeruginosa
Rhinovirus
Rubella virus
Salmonella
Salmonella typhi
Salmonella typhimurium
Shigella
Signal transduction, biological
Staphylococcus
Streptococcus
T cell (lymphocyte)
Treponema pallidum
Trichomonas vaginalis
Tritrichomonas foetus
Trypanosoma
Vaccines
Vibrio cholerae
Yersinia enterocolitica
Yersinia pestis
Yersinia pseudotuberculosis
   (vaccine comprising microbial adhesion mol. antigen as therapeutic and
     ***diagnostic***
                       for microbial infections)
Cell adhesion molecules
RL: ADV (Adverse effect, including toxicity); BSU (Biological study,
unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
   (vaccine comprising microbial adhesion mol. antigen as therapeutic and
     ***diagnostic***
                       for microbial infections)
Antiqens
Carbohydrates, biological studies
Glycolipids
Glycoproteins, general, biological studies
Integrins
Selectins
RL: BSU (Biological study, unclassified); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)
   (vaccine comprising microbial adhesion mol. antigen as therapeutic and
     ***diagnostic*** for microbial infections)
ADP ribosylation factor
Adhesins
Antiserums
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ΤТ

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ΤТ

Chemokines

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Cytokines
    Glycoconjugates
     Immunoglobulins
     LFA-1 (antigen)
    Ras proteins
    Rho protein (G protein)
     Toxins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (vaccine comprising microbial adhesion mol. antigen as therapeutic and
          ***diagnostic*** for microbial infections)
ΙT
    Cell adhesion molecules
    RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (vascular or VCAM; vaccine comprising microbial adhesion mol. antigen
       as therapeutic and ***diagnostic*** for microbial infections)
ΤТ
    Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.v; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΤТ
    Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.1.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΙT
     Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.2.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΙT
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.3.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΙT
     Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.4.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and
                        ***diagnostic*** for microbial infections)
ΙT
     Integrins
    RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.5.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and ***diagnostic*** for microbial infections)
ΙT
    Integrins
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (.alpha.6.beta.1; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and
                        ***diagnostic*** for microbial infections)
    72146-52-2, Mutan
ΤТ
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (Streptococcal; vaccine comprising microbial adhesion mol. antigen as
       therapeutic and
                         ***diagnostic*** for microbial infections)
ΤТ
     9005-32-7, Alginic acid
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (gel; vaccine comprising microbial adhesion mol. antigen as therapeutic
        and ***diagnostic*** for microbial infections)
ΙT
     59-23-4, Galactose, biological studies 63-42-3, Lactose
                                                                131-48-6,
    N-Acetylneuraminic acid
                              1811-31-0, N-Acetylgalactosamine
                                                                2438-80-4,
    Fucose 3416-24-8, Glucosamine 3458-28-4, Mannose 7512-17-6
     7535-00-4, Galactosamine
    RL: BSU (Biological study, unclassified); BIOL (Biological study)
```

(microbial adhesion mol. contq.; vaccine comprising microbial adhesion

- mol. antigen as therapeutic and ***diagnostic*** for microbial
 infections)
- IT 29350-58-1, PNAd-1
 - RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses) (vaccine comprising microbial adhesion mol. antigen as therapeutic and ***diagnostic*** for microbial infections)
- L11 ANSWER 3 OF 5 EMBASE COPYRIGHT (c) 2010 Elsevier B.V. All rights reserved on STN
- AN 2007510106 EMBASE <<LOGINID::20100824>>
- TI Rapid ***liposomal*** agglutination card test for the detection of antigens in patients with active ***tuberculosis*** .
- AU Tiwari, R.P.
- CS Diagnostic Division, Nicholas Piramal India Limited, Pawane, Navi, Mumbai, India.
- AU Tiwari, R.P.; Garg, S.K.; Bisen, Prakash S. (correspondence)
- CS Institute of Biotechnology and Allied Sciences, Seedling Academy of Design, Technology and Management, Jagatpura, Jaipur, India. psbisen@gmail.com
- AU Garg, S.K.
- CS Department of Biochemistry, University of Nebraska, Lincoln, NE, United States.
- AU Bharmal, R.N.; Kartikeyan, S.
- CS Department of Microbiology, Preventive and Social Medicine, Rajiv Gandhi Medical College, Kalwa, Thane, India.
- AU Bisen, Prakash S. (correspondence)
- CS Bisen Biotech and Biopharma Pvt. Ltd., M-7 Laxmipuram, Transport Nagar, Gwalior 474009, India. psbisen@gmail.com
- SO International Journal of Tuberculosis and Lung Disease, (Oct 2007) Vol. 11, No. 10, pp. 1143-1151.

 Refs: 30
 - ISSN: 1027-3719 CODEN: IJTDFO
- CY France
- DT Journal; Article
- FS 015 Chest Diseases, Thoracic Surgery and Tuberculosis 004 Microbiology: Bacteriology, Mycology, Parasitology and Virology 006 Internal Medicine
- LA English
- SL English; French; Spanish; Castilian
- ED Entered STN: 30 Oct 2007
 Last Updated on STN: 30 Oct 2007
- SETTING: A total of 1360 subjects with clinically confirmed pulmonary and AΒ ***tuberculosis*** (TB) and other non-tuberculous extra-pulmonary conditions. OBJECTIVES: To develop a rapid, sensitive and specific ***diagnostic*** test for the detection of the ***qlycolipid*** antigen of Mycobacterium ***tuberculosis*** in a variety of clinical samples. STUDY DESIGN: Affinity-purified rabbit anti- ***qlycolipid*** (IgG) were coupled to ***liposome*** particles ***antibodies*** (0.2-0.4 .mu.m) in the presence of 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride and N-hydroxysuccinamide to prepare the working reagent of the TB/M card test. RESULTS: ***Antibody*** -conjugated ***liposomes*** , when determined with the ***glycolipid*** antigens present in the specimens, formed a dark blue agglutination within 4 min. No dumping was observed in samples from normal healthy subjects or patients with other diseases. The test was shown to be effective in detecting ***glycolipid*** antigens of M. ***tuberculosis*** in clinical samples from patients with active TB with as low as 1 ng/ml

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analytical sensitivity, 97.4% clinical sensitivity and 96.9% specificity.
     CONCLUSION: The TB/M card test was found to be comparatively economical (4
     Indian Rupees or US$ 0.09/test), rapid (4 min) and seems fairly useful for
    mass testing of a variety of biological specimens (cerebrospinal, pleural
     and synovial fluids, serum, tissue biopsy extract) from patients with
     tuberculous meningitis, pulmonary TB and other extra-pulmonary TB in
     endemic countries. . COPYRGT. 2007 The Union.
TΤ
    Rapid ***liposomal*** agglutination card test for the detection of
     antigens in patients with active ***tuberculosis***
AΒ
     SETTING: A total of 1360 subjects with clinically confirmed pulmonary and
     extra-pulmonary ***tuberculosis*** (TB) and other non-tuberculous
     conditions. OBJECTIVES: To develop a rapid, sensitive and specific
      ***diagnostic*** test for the detection of the ***qlycolipid***
     antigen of Mycobacterium
                              ***tuberculosis*** in a variety of clinical
     samples. STUDY DESIGN: Affinity-purified rabbit anti- ***glycolipid***
       ***antibodies***
                         (IgG) were coupled to ***liposome***
     (0.2-0.4 .mu.m) in the presence of 1-ethyl-3-(3-dimethylaminopropyl)
     carbodiimide hydrochloride and N-hydroxysuccinamide to prepare the working
     reagent of the TB/M card test. RESULTS: ***Antibody*** -conjugated
       present in the specimens, formed a dark blue agglutination within 4 min.
    No dumping was observed in samples from normal healthy subjects or
    patients with other diseases. The test was shown to be effective in detecting ***glycolipid*** antigens of M. ***tuberculosis*** in
     clinical samples from patients with active TB with as low as 1 ng/ml
     analytical sensitivity, 97.4% clinical sensitivity and. . .
CT
    Medical Descriptors:
    adolescent
     adult
     *agglutination test
     *antigen detection
     article
     cerebrospinal fluid
     controlled study
         ***diagnostic test***
        ***extrapulmonary tuberculosis***
         ***lung tuberculosis***
    major clinical study
        ***Mycobacterium tuberculosis***
    pleura fluid
    priority journal
     school child
     sensitivity and specificity
     synovial fluid
         ****tuberculosis***
     tuberculous meningitis
     1 (3 dimethylaminopropyl) 3 ethylcarbodiimide
     amide
         ***antibody conjugate***
         ***qlycolipid***
         ***liposome***
     n hydroxysuccinamide
    tissue extract
L11 ANSWER 4 OF 5 EMBASE COPYRIGHT (c) 2010 Elsevier B.V. All rights
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reserved on STN

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0006205491 EMBASE <<LOGINID::20100824>>
ΑN
    MEDLINE.RTM. is the source for the citation and abstract of this record.
CP
ΤI
     [Major trends in lipid immunochemistry].
    Osnovnye napravleniia immunokhimii lipidov..
ΑU
    Shvets, V.I. (correspondence); Krasnopol'skii, I.M.
SO
    Ukrainskii biokhimicheskii zhurnal, (1984 May-Jun) Vol. 56, No. 3, pp.
     254-263.
    ISSN: 0201-8470
CY
    Russian Federation
DT
    Journal; Article
FS
    MEDLINE
LA
    Russian
ΕD
    Entered STN: Mar 2010
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AΒ
    Data are presented on immunochemical properties of lipids, the most
     important group of biologically active substances. Problems on antigenic,
     immunogenic and adjuvant activities of lipids are considered. A possible
     use of lipid antigens for
                                ***diagnosis*** of different infectious
     diseases is demonstrated and main principles of their construction are
     suggested. Data are available on immunogenicity of phospho- and
      ***glycolipid*** mixtures as well as on practical application of the
     obtained ***antibodies*** . Guidelines for the use of immunochemical
    properties of lipids are outlined.
     . . active substances. Problems on antigenic, immunogenic and
AB
     adjuvant activities of lipids are considered. A possible use of lipid
     antigens for ***diagnosis*** of different infectious diseases is
     demonstrated and main principles of their construction are suggested.
     Data are available on immunogenicity of phospho- and
                                                          ***alvcolipid***
    mixtures as well as on practical application of the obtained
      ***antibodies*** . Guidelines for the use of immunochemical properties
     of lipids are outlined.
    Medical Descriptors:
CT
    animal
     article
    brain
    cattle
    human
     immunization
     immunology
        ***lung tuberculosis: DI, diagnosis***
         ***schistosomiasis: DI, diagnosis***
     serology
     syphilis serology
     cardiolipin
         ***diagnostic agent***
     *epitope: AN, drug analysis
     immunological adjuvant: AD, drug administration
     *lipid: AD, drug administration
         ***liposome: AD, drug administration***
L11 ANSWER 5 OF 5 SCISEARCH COPYRIGHT (c) 2010 The Thomson Corporation on
     STN
     2008:599597 SCISEARCH <<LOGINID::20100824>>
ΑN
GΑ
    The Genuine Article (R) Number: 292ZK
ΤI
    A novel application of affinity biosensor technology to detect
       ***antibodies*** to mycolic acid in ***tuberculosis*** patients
ΑU
    Verschoor, Jan A. (Reprint)
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CS
    Univ Pretoria, Dept Biochem, ZA-0002 Pretoria, South Africa (Reprint)
     Thanyani, Simon T.; Roberts, Vanessa; Siko, D. Gilbert R.; Vrey, Pieter
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    E-mail: jan.verschoor@up.ac.za
CYA South Africa
    JOURNAL OF IMMUNOLOGICAL METHODS, (20 MAR 2008) Vol. 332, No. 1-2, pp.
SO
     61 - 72.
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PB
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    Article; Journal
LA
    English
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ED
    Entered STN: 15 May 2008
     Last Updated on STN: 15 May 2008
     *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
AΒ
          ***Tuberculosis*** has re-emerged as a global health problem due to
     co-infection with HIV and the emergence of drug-resistant strains of
    Mycobacterium ***tuberculosis*** . HIV co-infection introduced a 30%
     underestimation in TB
                           ***diagnosis*** based on sputum analysis,
     calling for a reliable and fast serodiagnostic assay to assist in the
     management of TB in HIV-burdened populations. Serodiagnosis with
    mycobacterial lipid cell wall antigens gave promising results, in
    particular with LAM and cord factor. Free mycolic acids have also been
     considered because they are unique in structure to each species of
    Mycobacterium and can be economically extracted and purified. In a
     standard immunoassay such as ELISA, however, an unacceptable number of
     false positive and false negative test results were obtained. Here we
     report a much improved biosensor method to detect ***antibodies*** to
    mycolic acids in patient serum as surrogate markers of active
       ***tuberculosis*** . Mycolic acid (MA)
                                                ***liposomes***
     immobilized on a non-derivatized twin-celled biosensor cuvette and blocked
     with saponin. A high dilution of serum was used to calibrate the binding
     signal of the two cells, followed by contact with patient serum at a
     lesser dilution, but pre-incubated with either antigen-carrying, or empty
       ***liposomes*** . The serum, or the protein A purified IgG thereof,
from
                     ***tuberculosis*** patients could be inhibited from
     sputum-positive
     binding to the MA in the biosensor by prior incubation with MA-containing
      ***liposomes*** . The accuracy of the inhibition test was 84% if
     HIV-positive patients for whom a negative TB sputum analyses could not be
     relied upon to serve as a reference standard were excluded. If biosensor
     technology could be made suitable for high throughput screening, then it
    may provide the solution to the serodiagnosis of ***tuberculosis***
     against a background of HIV (C) 2007 Elsevier B.V. All rights reserved.
ΤI
    A novel application of affinity biosensor technology to detect
       ***antibodies***
                        to mycolic acid in ***tuberculosis***
                                                                  patients
AB
          ***Tuberculosis***
                             has re-emerged as a global health problem due to
     co-infection with HIV and the emergence of drug-resistant strains of
     Mycobacterium ***tuberculosis*** . HIV co-infection introduced a 30%
     underestimation in TB ***diagnosis*** based on sputum analysis,
     calling for a reliable and fast serodiagnostic assay to assist in the
    management of TB in. . . of false positive and false negative test
    results were obtained. Here we report a much improved biosensor method to
    detect ***antibodies*** to mycolic acids in patient serum as surrogate
    markers of active ***tuberculosis*** . Mycolic acid (MA)
       ***liposomes*** were immobilized on a non-derivatized twin-celled
    biosensor cuvette and blocked with saponin. A high dilution of serum was
     used to. . . the two cells, followed by contact with patient serum at a
```

lesser dilution, but pre-incubated with either antigen-carrying, or empty ***liposomes*** . The serum, or the protein A purified IgG thereof,

from

sputum-positive ***tuberculosis*** patients could be inhibited from binding to the MA in the biosensor by prior incubation with MA-containing ***liposomes***. The accuracy of the inhibition test was 84% if HIV-positive patients for whom a negative TB sputum analyses could not. . biosensor technology could be made suitable for high throughput screening, then it may provide the solution to the serodiagnosis of ***tuberculosis*** against a background of HIV (C) 2007 Elsevier B.V. All rights reserved.

- ST Author Keywords: ***antibodies*** ; mycolic acids; biosensor; Mycobacterium ***tuberculosis*** ; serodiagnosis
- STP KeyWords Plus (R): MYCOBACTERIUM- ***TUBERCULOSIS*** ; PULMONARY

 TUBERCULOSIS ; IMMUNE-RESPONSES; PROTEIN-A; BIOMOLECULAR

 INTERACTIONS; SEROLOGICAL ***DIAGNOSIS*** ; ***GLYCOLIPID***

 ANTIGEN; OPTICAL BIOSENSOR; RESONANT MIRROR; IGG ***ANTIBODY***